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NEWS	3	MAR 31	CAS REGISTRY enhanced with additional experimental spectra
NEWS	4	MAR 31	CA/CAPLUS and CASREACT patent number format for U.S. applications updated
NEWS	5	MAR 31	LPCI now available as a replacement to LDPCI
NEWS	6	MAR 31	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS	7	APR 04	STN AnaVist, Version 1, to be discontinued
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NEWS	12	MAY 30	DGENE, PCTGEN, and USGENE enhanced with new homology sequence search option
NEWS	13	JUN 06	EPFULL enhanced with 260,000 English abstracts
NEWS	14	JUN 06	KOREAPAT updated with 41,000 documents
NEWS	15	JUN 13	USPATFULL and USPAT2 updated with 11-character patent numbers for U.S. applications
NEWS	16	JUN 19	CAS REGISTRY includes selected substances from web-based collections
NEWS	17	JUN 25	CA/CAPLUS and USPAT databases updated with IPC reclassification data
NEWS	18	JUN 30	AEROSPACE enhanced with more than 1 million U.S. patent records
NEWS	19	JUN 30	EMBASE, EMBAL, and LEMBASE updated with additional options to display authors and affiliated organizations
NEWS	20	JUN 30	STN on the Web enhanced with new STN AnaVist Assistant and BLAST plug-in
NEWS	21	JUN 30	STN AnaVist enhanced with database content from EPFULL
NEWS	22	JUL 28	CA/CAPLUS patent coverage enhanced
NEWS	23	JUL 28	EPFULL enhanced with additional legal status information from the EPOLINE Register
NEWS	24	JUL 28	IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
NEWS	25	JUL 28	STN Viewer performance improved
NEWS	26	AUG 01	INPADOCDB and INPAFAMDB coverage enhanced
NEWS	27	AUG 13	CA/CAPLUS enhanced with printed Chemical Abstracts page images from 1967-1998
NEWS	28	AUG 15	CAOLD to be discontinued on December 31, 2008
NEWS	29	AUG 15	CAPLUS currency for Korean patents enhanced
NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3,			

AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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=> s p2x2
L1 2066 P2X2

=> s l1 and antisense
L2 35 L1 AND ANTISENSE

=> dup rem l2
PROCESSING COMPLETED FOR L2
L3 23 DUP REM L2 (12 DUPLICATES REMOVED)

=> d ti 1-23

L3 ANSWER 1 OF 23 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on STN
TI P2X(3) receptors and sensory transduction.

L3 ANSWER 2 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 1
TI Painful purinergic receptors

L3 ANSWER 3 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI P2X3 receptors and sensory transduction

L3 ANSWER 4 OF 23 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights reserved on STN

TI Ion channels in pain transmission.

L3 ANSWER 5 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI The peripheral modulation of nociceptive or pain signals mediated by P2X receptors

L3 ANSWER 6 OF 23 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights reserved on STN

TI Pharmacology of P2X channels.

L3 ANSWER 7 OF 23 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights reserved on STN

TI Purinergic control of neuropathic pain.

L3 ANSWER 8 OF 23 MEDLINE on STN

TI Inhibitory role of supraspinal P2X3/P2X2/3 subtypes on nociception in rats.

L3 ANSWER 9 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 2

TI Inhibitory role of supraspinal P2X3/P2X2/3 subtypes on nociception in rats

L3 ANSWER 10 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI P2X receptors: targets for novel analgesics?

L3 ANSWER 11 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI Treatment of neurological disorders by double-stranded RNA (dsRNA) administration

L3 ANSWER 12 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 3

TI P2X3 receptors and peripheral pain mechanisms

L3 ANSWER 13 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI Experimental investigation of P2 receptors and pain

L3 ANSWER 14 OF 23 MEDLINE on STN DUPLICATE 4

TI Chronic pain and microglia: the role of ATP.

L3 ANSWER 15 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI Method for identifying compounds to treat urological disorders by measuring their effect on disease-related genes and proteins, and use of the identified compounds in treatment

L3 ANSWER 16 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI Antisense oligonucleotides regulating rat and human purinoreceptor P2X3 gene expression and their use in pain therapy

L3 ANSWER 17 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

TI Contributions of P2X3 homomeric and heteromeric channels to acute and chronic pain

L3 ANSWER 18 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 5

TI ATP induced three types of pain behaviors, including allodynia

L3 ANSWER 19 OF 23 MEDLINE on STN DUPLICATE 6

TI Downregulation of P2X3 receptor-dependent sensory functions in A/J inbred mouse strain.

L3 ANSWER 20 OF 23 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on
STN
TI ATP-induced three types of pain behaviors including allodynia.

L3 ANSWER 21 OF 23 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on
STN
TI Peripheral and spinal mechanisms of alpha,beta-methylene ATP-induced
mechanical allodynia in rats.

L3 ANSWER 22 OF 23 MEDLINE on STN
TI P2 receptors in the thymus: expression of P2X and P2Y receptors in adult
rats, an immunohistochemical and in situ hybridisation study.

L3 ANSWER 23 OF 23 MEDLINE on STN DUPLICATE 7
TI Phosphorothioate oligodeoxynucleotides can selectively alter neuronal
activity in the cochlea.

=> d ab 16 22 23

L3 ANSWER 16 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
AB The invention relates to antisense oligonucleotides, compns. and
methods useful for modulating the expression of P2X2. The
compns. comprise antisense oligonucleotides targeted to nucleic
acids encoding P2X2.

L3 ANSWER 22 OF 23 MEDLINE on STN
AB The expression of the seven P2X receptor subtypes and of two P2Y receptors
was examined immunohistochemically and by in situ hybridisation in thymi
of adult male rats. P2X4, P2Y2 and 4 receptor mRNA colocalisation studies
combining in situ hybridisation and immunohistochemistry were also carried
out. P2X and P2Y receptors were found on thymocytes. P2X receptors were
also abundant in cells of the thymic microenvironment, involved in control
of T-cell maturation in vivo. We are the first to describe the expression
of P2X4 receptors on thymocytes and confirm the finding of P2X1 and P2Y2
receptors on subpopulations of lymphocytes. P2X1,2,3,4 and 5 receptors
were present in blood vessels of the thymus. P2X1,2 and 4 receptors were
detected in vascular smooth muscle, while P2X3 receptors appeared to be
associated with endothelial cells; some small arteries were positive for
P2X5, possibly labelling vascular smooth muscle or fibroblasts in the
adventitia. P2X2,3,6 and 7 receptors were found on thymic
epithelial cells. P2X2 and 3 receptors were abundant on
medullary epithelial cells, whilst P2X6 receptors were prominent in
Hassall's corpuscles. P2X2 receptors were found on subcapsular
and perivascular epithelial cells. P2X2,6 and 7 receptors were
detected in epithelial cells along the thymic septa. Expression of P2X
receptors was also investigated by Western blotting of crude thymic tissue
extracts under reducing conditions. All seven P2X receptor subtypes were
found to be dimers of approximately 70 kDa and 140 kDa molecular weight.
ATP-mediated apoptosis and cell proliferation of thymocytes are discussed.

L3 ANSWER 23 OF 23 MEDLINE on STN DUPLICATE 7
AB A growing body of evidence indicates that extracellular adenosine
triphosphate (ATP) may have a major role in cochlear function.
Antagonists of ionotropic ATP receptors (P2X2) have significant
effects on cochlear potentials and distortion product otoacoustic
emissions (DPOAEs). We tested whether antisense
oligodeoxynucleotides (ODNs) would mimic the functional deficiencies
induced by the ATP antagonists through binding to P2X2 ATP
receptor mRNA and thereby reduce the number of ATP receptors expressed in
the membrane of the cells. Both a phosphorothioate ODN (S-ODN)
antisense and a phosphodiester ODN (P-ODN) antisense to

the P2X2 sequence and random sense ODNs containing 21 nucleotides were administered chronically (7 days) to the guinea pig cochlea via the perilymph compartment. Sound evoked cochlear potentials (cochlear microphonic; summating potential; compound action potential of the auditory nerve, CAP; latency of the first negative peak in the CAP, N1 latency) and DPOAEs were monitored to assess the effects of the ODNs. Results indicate that the phosphorothioate derivatives of both the antisense and random sense ODNs suppressed the CAP and prolonged the N1 latency with no significant effect on the other parameters. The P-ODNs had no effect. Since both the antisense and random sense S-ODNs had the same effect, we conclude that the S-ODNs affected neuronal function in a manner that did not involve binding to the ATP receptor mRNA.

=> d 16 23

L3 ANSWER 16 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2003:551668 CAPLUS

DN 139:95485

TI Antisense oligonucleotides regulating rat and human purinoreceptor P2X3 gene expression and their use in pain therapy

IN Shuster, Samuel J.; Arvidsson, Ulf N. G.; Stone, Laura S.; Zhang, Hong-yan; Hart, Lucy Vulchanova

PA Algos Therapeutics, Inc., USA

SO PCT Int. Appl., 35 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003057898	A2	20030717	WO 2002-US41833	20021231
	WO 2003057898	A3	20031023		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	AU 2002360847	A1	20030724	AU 2002-360847	20021231
	EP 1470145	A2	20041027	EP 2002-796136	20021231
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK			
	JP 2005514034	T	20050519	JP 2003-558191	20021231
	US 20080119423	A1	20080522		
PRAI	US 2001-346155P	P	20011231	US 2004-500449	20041203
	WO 2002-US41833	W	20021231		

L3 ANSWER 23 OF 23 MEDLINE on STN

DUPLICATE 7

AN 1999419708 MEDLINE

DN PubMed ID: 10491959

TI Phosphorothioate oligodeoxynucleotides can selectively alter neuronal activity in the cochlea.

AU LeBlanc C S; Fallon M; Parker M S; Skellett R; Bobbin R P

CS Department of Otorhinolaryngology and Biocommunication, Louisiana State University Medical Center, New Orleans 70112-2234, USA.

NC T32-DC00007 (United States NIDCD)

SO Hearing research, (1999 Sep) Vol. 135, No. 1-2, pp. 105-12.
Journal code: 7900445. ISSN: 0378-5955.
CY Netherlands
DT Journal; Article; (JOURNAL ARTICLE)
(RESEARCH SUPPORT, NON-U.S. GOV'T)
(RESEARCH SUPPORT, U.S. GOV'T, P.H.S.)
LA English
FS Priority Journals
EM 199911
ED Entered STN: 11 Jan 2000
Last Updated on STN: 11 Jan 2000
Entered Medline: 5 Nov 1999

=> d ab 15 19

L3 ANSWER 15 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN
AB The present invention relates to methods for the diagnosis and treatment of a urol. disorder or urol. disorders. Specifically, the present invention identifies the differential expression of 1435, 559, 34021, 44099, 25278, 641, 260, 55089, 21407, 42032, 46656, 62553, 302, 323, 12303, 985, 13237, 13601, 18926, 318, 2058 and 6351 genes in tissues relating to urol. disorders, relative to their expression in normal, or non-urol. disorders, and/or in response to manipulations relevant to a urol. disorder. The present invention describes methods for the diagnostic evaluation and prognosis of various urol. disorders, and for the identification of subjects exhibiting a predisposition to such conditions. glucose. The invention also provides methods for identifying a compound capable of modulating a urol. disorder or urol. disorders. The present invention also provides methods for the identification and therapeutic use of compds. as treatments of urol. disorders.

L3 ANSWER 19 OF 23 MEDLINE on STN DUPLICATE 6
AB There is large variability in the various pain responses including those to tissue injury among inbred mouse strains. However, the determinant factors for the strain-specific differences remain unknown. The P2X3 sensory-specific ATP-gated channel has been implicated as a damage-sensing molecule that evokes a pain sensation by receiving endogenous ATP from injured tissue. In this study, to clarify the contribution of the sensory P2X3 signalling to strain-specific differences in tissue injury pain, we examined whether the P2X3-mediated in vivo and in vitro responses in dorsal root ganglion (DRG) neurons are changed in the A/J inbred mouse strain, which is known to be resistant to tissue injury pain caused by formalin. Here we found that A/J mice exhibited a low magnitude of nocifensive behaviour induced by the P2X agonist alpha,beta-methylene ATP (alpha beta meATP) into the hindpaw compared with C57BL/6 J mice. This behaviour was blocked by P2X3 antisense oligodeoxynucleotides. The low magnitude of the in vivo pain sensation could be observed similarly in the in vitro response; the increase in the intracellular Ca(2+) increase by alpha beta meATP in capsaicin-sensitive DRG neurons from A/J mice was significantly lower than that from C57BL/6 J mice. In A/J DRG neurons the P2X3 protein level was significantly lower compared with C57BL/6 J DRG neurons. The change in P2X3 protein was selective because P2X2 protein was expressed equally in both strains. The present study suggests that the downregulation of sensory P2X3 could be one of the molecular predispositions to low sensitivity to tissue injury pain in the A/J inbred mouse strain.

=> d 15 19

L3 ANSWER 15 OF 23 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2003:590937 CAPLUS
 DN 139:143886
 TI Method for identifying compounds to treat urological disorders by measuring their effect on disease-related genes and proteins, and use of the identified compounds in treatment
 IN Silos-Santiago, Inmaculada; Karicheti, Venkateswarlu
 PA Millennium Pharmaceuticals, Inc., USA
 SO PCT Int. Appl., 242 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003061573	A2	20030731	WO 2003-US1450	20030116
	WO 2003061573	A3	20031231		
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
	US 20030148394	A1	20030807	US 2003-345680	20030116
	EP 1472376	A2	20041103	EP 2003-731962	20030116
	R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK			
	JP 2005514940	T	20050526	JP 2003-561519	20030116
	US 20060088881	A1	20060427	US 2005-302678	20051214
PRAI	US 2002-349511P	P	20020118		
	US 2002-360500P	P	20020228		
	US 2002-365041P	P	20020315		
	US 2002-374063P	P	20020419		
	US 2002-403468P	P	20020814		
	US 2002-414262P	P	20020927		
	US 2002-419986P	P	20021021		
	US 2002-423809P	P	20021105		
	US 2002-429797P	P	20021126		
	US 2003-345680	B1	20030116		
	WO 2003-US1450	W	20030116		

L3 ANSWER 19 OF 23 MEDLINE on STN DUPLICATE 6
 AN 2002323765 MEDLINE
 DN PubMed ID: 12028354
 TI Downregulation of P2X3 receptor-dependent sensory functions in A/J inbred mouse strain.
 AU Tsuda Makoto; Shigemoto-Mogami Yukari; Ueno Shinya; Koizumi Schuichi; Ueda Hiroshi; Iwanaga Toshihiko; Inoue Kazuhide
 CS Section of Neuroparmacology, Division of Pharmacology, National Institute of Health Sciences, Tokyo, Japan.
 SO The European journal of neuroscience, (2002 May) Vol. 15, No. 9, pp. 1444-50.
 Journal code: 8918110. ISSN: 0953-816X.
 CY France
 DT (COMPARATIVE STUDY)
 Journal; Article; (JOURNAL ARTICLE)
 (RESEARCH SUPPORT, NON-U.S. GOV'T)
 LA English
 FS Priority Journals

EM 200207
ED Entered STN: 18 Jun 2002
Last Updated on STN: 27 Jul 2002
Entered Medline: 26 Jul 2002

=> s p2rx2
L4 25 P2RX2

=> s l4 and antisense
L5 0 L4 AND ANTISENSE

=> s purinergic receptor p2x
L6 148 PURINERGIC RECEPTOR P2X

=> s l6 and antisense
L7 3 L6 AND ANTISENSE

=> s l7 not l3
L8 3 L7 NOT L3

=> dup rem l8
PROCESSING COMPLETED FOR L8
L9 1 DUP REM L8 (2 DUPLICATES REMOVED)

=> d l

L9 ANSWER 1 OF 1 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights reserved on STN DUPLICATE 1
AN 2005310641 EMBASE
TI Mechanical strain opens connexin 43 hemichannels in osteocytes: A novel mechanism for the release of prostaglandin.
AU Cherian, Priscilla P.; Siller-Jackson, Arlene J.; Gu, Sumin; Wang, Xin; Jiang, Jean X. (correspondence)
CS Department of Biochemistry, University of Texas Health Science Center, San Antonio, TX 78229-3900, United States. jiangj@uthscsa.edu
AU Sprague, Eugene
CS Department of Radiology, University of Texas Health Science Center, San Antonio, TX 78229-3900, United States.
AU Bonewald, Lynda F.
CS Department of Oral Biology, School of Dentistry, University of Missouri, Kansas City, MO 64108, United States.
SO Molecular Biology of the Cell, (Jul 2005) Vol. 16, No. 7, pp. 3100-3106.
Refs: 39
ISSN: 1059-1524 CODEN: MBCEEV
CY United States
DT Journal; Article
FS 029 Clinical and Experimental Biochemistry
LA English
SL English
ED Entered STN: 5 Aug 2005
Last Updated on STN: 5 Aug 2005

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TOTAL

ENTRY

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